



上海源叶生物科技有限公司
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产品名称: ASK1-IN-1

产品别名: GS-444217

生物活性:

Description	GS-444217 is a potent, orally available and selective ATP-competitive inhibitor of apoptosis signal-regulating kinase 1 (ASK1) with an IC50 of 2.87 nM[1].																				
IC ₅₀ & Target	ASK1 2.87 nM (IC ₅₀)																				
In Vitro	Treatment with GS-444217 reduces ASK1 phosphorylation and prevents the phosphorylation of MKK3/6, MKK4, p38, and JNK at concentrations of 0.3 μM and above with full suppression of ASK1 activity at 1 μM. GS-444217 (1 μM) reduces ASK1 activity within 5 minutes of addition to the cultures, reaching a maximum level of inhibition by 30 minutes. Removal of GS-444217 from the cultures results in reactivation of ASK1 autoprophosphorylation within 10 minutes and near-complete recovery 2 hours after drug washout ^[1] .																				
In Vivo	GS-444217 reduces oxidative stress (OS)-induced ASK1 signaling in kidney and inhibits acute renal tubular injury in rats. GS-444217 (30 mg/kg) inhibits activation of ASK1, p38, and JNK in rat kidney. GS-444217 has an in vivo EC50 of approximately 1.6 μM for inhibiting the ASK1 pathway in rodent kidney[1].																				
Solvent&Solubility	<p>In Vitro: DMSO : 15.5 mg/mL (37.67 mM; Need ultrasonic and warming)</p> <table border="1"><thead><tr><th rowspan="2">Preparing Stock Solutions</th><th>Solvent / Mass Concentration</th><th>1 mg</th><th>5 mg</th><th>10 mg</th></tr></thead><tbody><tr><td>1 mM</td><td>2.4304 mL</td><td>12.1518 mL</td><td>24.3037 mL</td></tr><tr><td>5 mM</td><td>0.4861 mL</td><td>2.4304 mL</td><td>4.8607 mL</td></tr><tr><td>10 mM</td><td>0.2430 mL</td><td>1.2152 mL</td><td>2.4304 mL</td></tr></tbody></table> <p>*请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液。一旦配成溶液,请分装保存,避免反复冻融造成的产品失效。 储备液的保存方式和期限 -80°C, 6 months; -20°C, 1 month。-80°C 储存时,请在 6 个月内使用, -20°C 储存时,请在 1 个月内使用。</p> <p>In Vivo: 请根据您的实验动物和给药方式选择适当的溶解方案。以下溶解方案都请先按照 In Vitro 方式配制澄清的储备液,再依次添加助溶剂: ——为保证实验结果的可靠性,澄清的储备液可以根据储存条件,适当保存;体内实验的工作液,建议您现用现配,当天使用;以下溶剂前显示的百分比是指该溶剂在您配制终溶液中的体积占比;如在配制过程中出现沉淀、析出现象,可以通过加热和/或超声的方式助溶 1.请依序添加每种溶剂: 10% DMSO→40% PEG300 →5% Tween-80 → 45% saline Solubility: ≥ 2.5 mg/mL (6.08 mM); Clear solution 此方案可获得 ≥ 2.5 mg/mL (6.08 mM, 饱和度未知) 的澄清溶液。 以 1 mL 工作液为例,取 100 μL 25.0 mg/mL 的澄清 DMSO 储备液加到 400 μL PEG300 中,混合均匀; 向上述体系中加入 50 μL Tween-80, 混合均匀;然后继续加入 450 μL 生理盐水定容至 1 mL。 2.请依序添加每种溶剂: 10% DMSO→ 90% (20% SBE-β-CD in saline)</p>				Preparing Stock Solutions	Solvent / Mass Concentration	1 mg	5 mg	10 mg	1 mM	2.4304 mL	12.1518 mL	24.3037 mL	5 mM	0.4861 mL	2.4304 mL	4.8607 mL	10 mM	0.2430 mL	1.2152 mL	2.4304 mL
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	<p>Solubility: $\geq 2.5 \text{ mg/mL}$ (6.08 mM); Clear solution 此方案可获得 $\geq 2.5 \text{ mg/mL}$ (6.08 mM, 饱和度未知) 的澄清溶液。 以 1 mL 工作液为例, 取 100 μL 25.0 mg/mL 的澄清 DMSO 储备液加到 900 μL 20% 的 SBE-β-CD 生理盐水水溶液中, 混合均匀。</p> <p>3. 请依序添加每种溶剂: 10% DMSO → 90% corn oil Solubility: $\geq 2.5 \text{ mg/mL}$ (6.08 mM); Clear solution 此方案可获得 $\geq 2.5 \text{ mg/mL}$ (6.08 mM, 饱和度未知) 的澄清溶液, 此方案不适用于实验周期在半个月以上的实验。 以 1 mL 工作液为例, 取 100 μL 25.0 mg/mL 的澄清 DMSO 储备液加到 900 μL 玉米油中, 混合均匀。</p>
References	<p>[1]. Liles JT, et al. ASK1 contributes to fibrosis and dysfunction in models of kidney disease. <i>J Clin Invest.</i> 2018 Oct 1;128(10):4485-4500. [2]. Budas GR, et al. ASK1 Inhibition Halts Disease Progression in Preclinical Models of Pulmonary Arterial Hypertension. <i>Am J Respir Crit Care Med.</i> 2018 Feb 1;197(3):373-385.</p>



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